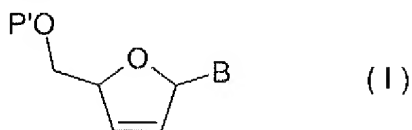


## AMENDMENTS TO THE CLAIMS

This listing of claims will replace all prior versions, and listings, of claims in the application:

### Listing of Claims:

1. (Currently Amended) A process for preparing 2',3'-dideoxynucleoside of formula

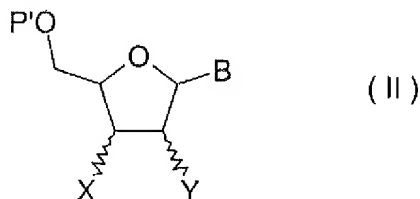


in which

P' represents hydrogen or a protecting group P, and

B represents a natural or modified on the ring by aza-substitution, optionally substituted fluorinated or methylated purine or pyrimidine base or a five- or six-membered monocyclic or eleven- or twelve-membered bicyclic, optionally substituted fluorinated or methylated heterocyclic system containing at least one nitrogen atom;

which comprises reducing a compound of formula



in which

X and Y represent, alternately, a halogen or an acyloxy group RCOO-,

P' and B have the meanings given above,

by reaction with divalent zinc and an activating agent in an organic phase to provide the compound of formula 1, and,

adding a sulfide solution of an alkali metal sulfide or alkaline-earth metal sulfide to precipitate divalent zinc as zinc sulfide from said organic phase.

2. (Previously Presented) The process according to Claim 1, in which:

- P' represents an acyl group RCO-, in which R represents a C<sub>1</sub>-C<sub>5</sub> alkyl or a group R<sup>1</sup>COOC(R<sup>2</sup>R<sup>3</sup>)-, in which R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> represent a C<sub>1</sub>-C<sub>5</sub> alkyl;
- B represents an optionally substituted natural purine or pyrimidine base;
- X and Y represent, alternatively, bromine and an acyloxy group RCOO-, in which R represents a C<sub>1</sub>-C<sub>5</sub> alkyl, or a group R<sup>1</sup>COOC(R<sup>2</sup>R<sup>3</sup>)-, in which R<sup>1</sup>, R<sup>2</sup> and R<sup>3</sup> represent a C<sub>1</sub>-C<sub>5</sub> alkyl.

3. (Previously Presented) The process according to Claim 1, in which the said activating agent is selected from the group consisting of copper, acetic acid, ammonium salt, phosphonium salt, and mixtures thereof.

4. (Previously Presented) The process according to Claim 1, in which the said organic phase is a solvent selected from the group consisting of tetrahydrofuran, dimethylacetamide, alcohol, acetonitrile, chlorinated solvent, dimethyl sulfoxide, and mixtures thereof.

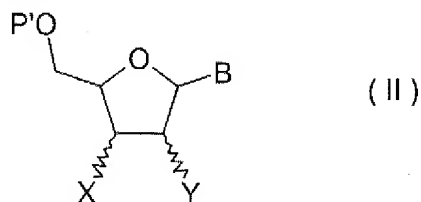
5. (Previously Presented) The process according to Claim 1, in which the said sulfide solution comprises a polar solvent selected from the group consisting of a dipolar aprotic solvent, water, and mixtures thereof.

6. (Previously Presented) The process according to Claim 1, in which said sulfide solution comprises the alkali metal sulfide or alkaline-earth metal sulfide in an amount of at least one molar equivalent relative to the starting material.

7. (Previously Presented) The process according to Claim 1, in which the alkali metal sulfide is sodium sulfide.

8. (Previously Presented) The process according to Claim 1, further comprising removing precipitated zinc sulfide by filtration.

9. (Previously Presented) The process according to Claim 1, which further comprises reducing the double bond of the compound of formula I to give the corresponding 2',3'-dideoxynucleoside of formula

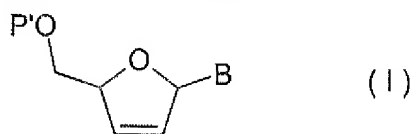


in which  $X = Y = H$ , and

P' represents an acyl group  $RCO-$ , in which R represents a  $C_1-C_5$  alkyl or a group  $R^1COOC(R^2R^3)-$ , in which  $R^1$ ,  $R^2$  and  $R^3$  represent a  $C_1-C_5$  alkyl;

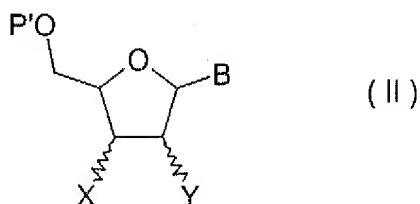
B represents an optionally substituted natural purine or pyrimidine base.

10. (Previously Presented) The process according to Claim 1, which further comprises the deprotection reaction of a compound of formula



in which P' represents a protecting group P, and B represents an optionally substituted natural purine or pyrimidine base, to give the corresponding compound of formula I, in which P' represents hydrogen.

11. (Previously Presented) Process according to Claim 9, which further comprises the deprotection reaction of a compound of formula



in which P' represents a protecting group P, X and Y represent H, and B represents an optionally substituted natural purine or pyrimidine base, to give the corresponding compound of formula II, in which P' represents hydrogen.

12. (Currently Amended) The process of claim 1, wherein the 2',3'-didehydro-2',3'-dideoxynucleoside is selected from the group consisting of 5-fluoro-2',3'-dideoxy-2',3'-didehydro- $\beta$ -D-cytidine, stavudine, dideoxyadenosine[[.]], didanosine, zalcitabine, and mixtures thereof.

13. (Currently Amended) The process of claim 1, wherein B is selected from the group consisting of adenine, inosine [[.]], 5-F-cytosine, hypoxanthine, thymine, and mixtures thereof.

14. (Previously Presented) The process of Claim 1, wherein said activating agent is selected from the group consisting of ammonium salt, phosphonium salt, and mixtures thereof.

15. (Previously Presented) The process of Claim 1, wherein said sulfide solution comprises the alkali metal sulfide or alkaline-earth metal sulfide in an amount greater than one molar equivalent relative to the divalent zinc.

16. (Previously Presented) The process of claim 2, wherein R<sup>1</sup> is methyl.

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17. (New) The process of claim 1, wherein said sulfide solution is at a temperature between 40 to 60 °C.